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⑮ Method for introducing foreign genes into green algae utilizing T-DNA of agrobacterium.

⑮ A genetically engineered green algal cell and a method for introducing biologically functional foreign genes into green algae are defined in which the T-DNA sequence of the *Agrobacterium* plasmid is used as a vector to transfer prokaryotic or eukaryotic DNA into the algal genome where it is replicated and expressed.

EP 0 108 580 A1

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- 1 -

METHOD FOR INTRODUCING FOREIGN GENES INTO
GREEN ALGAE UTILIZING T-DNA OF AGROBACTERIUM

5 This invention relates to a method for replica-
tion and expression of exogenous genes in algae.
More particularly, this invention relates to the
use of Agrobacterium as a vector in the genetic
engineering of green algae.

10 Algae are a very large group of photosynthetic
organisms. The group includes single celled and
multi-cellular organisms. There are green, blue-
green, red, and brown algae. The organisms have
flexible growth requirements and can grow in either
15 marine or fresh water environments. All algae are
able to use carbon dioxide and light as their carbon
and energy sources. They are among the most efficient
organisms to convert solar energy into organic matter.
Certain species have measured photosynthetic ef-
ficiencies of 1 to 2 per cent in the field, which
20 represents a biomass production of 15-30 g dry
wt/m²/day. Goldman, J. G. In: Algae Biomass eds.
G. Shelef and C. J. Soeder, 1980, Elsevier/North
Holland Biomedical Press, pp. 343-359. Under optimal
laboratory conditions, photosynthetic efficiencies
25 of 20 per cent have been reported. Shelef, G.;
Oswald, W. J.; and McGauhey, P. H. J. Sanit. Engin.
Div. Am. Soc. Civ. Engrs., vol. 96, SA1 (1970), pp.
91-110.

30 Algae have a varied biosynthetic machinery, and
many species produce compounds that have future
industrial applications. Some of these chemicals
are produced by algae in very high levels. One
species synthesizes over 50 per cent of its cell
mass as glycerol, and others produce high levels of
35 hydrocarbons, lipids, and protein. In addition, a
number of unique compounds are made by algae and

11 12 11 90

0108580

-2-

have potential applications in many different areas.
Aaronson, S.; Berner, T.; and Dubinsky, Z. In: Algae
Biomass, supra, pp. 575-601.

5 It is the biosynthetic capability of algae which
makes them a future source for valuable compounds.
Characteristics that make algae attractive for in-
dustrial purposes include their ability to use simple
components such as carbon dioxide and light for
growth, and their varied and efficient biosynthetic
10 capability.

The ultimate objective of the genetic engineer-
ing process is the transfer of a foreign gene coding
for a particular useful characteristic into a new
host cell where the foreign gene is then expressed.
15 The transfer of a foreign gene involves several
steps. First, the foreign gene coding for the
desired characteristic is incorporated into a vector
through a gene-splicing procedure. Vectors, or
cloning vehicles, are DNA (deoxyribonucleic acid)
20 molecules that physically transport the foreign
gene to the new host cell. Vectors are generally
either plasmids, which are circular DNA molecules,
or viruses, which are linear DNA molecules. The
vector-foreign gene hybrid is then transferred to
25 the host cell using a transformation process. The
result is a genetically engineered cell with the
desired foreign gene now present in the new host
cell.

A successfully genetically engineered cell is
30 one in which the foreign gene, and hence the desir-
ed characteristic, is "expressed." Gene expression
is a multistep process requiring transcription of
DNA into mRNA (messenger ribonucleic acid) and trans-
lation of mRNA into protein. For these purposes,
35 the DNA is divided into control regions, which
mediate expression of the information, and structural

M 12.11.90

0108580

-3-

regions, which code for proteins. To begin transcrip-
tion, the host cell RNA-polymerase must "recognize,"
or bind to, a control region of the DNA (hereinafter
referred to as the "promoter"). The promoter se-
5 quence is often different depending upon the DNA
material used, e.g., prokaryotic DNA v. eukaryotic
DNA. The enzyme RNA polymerase moves along the
structural gene transcribing the DNA encoded message
into mRNA. The mRNA must contain another control
10 sequence, the translation start site, which allows
it to bind to the ribosomes where it is translated
into protein. Several attempts to create genetic
engineering systems have failed because although
foreign DNA was successfully transferred to a new
15 host cell, the cell's own transcription and transla-
tion systems were unable to recognize the control
sequences of the foreign DNA and the foreign gene
was not expressed. Hopwood, David A., "The Genetic
Programming of Industrial Microorganisms," Scientific
20 American, vol. 245, No. 3, pp. 90-102, September
1981.

There are presently only a few species in which
the various steps for genetic engineering have been
determined. For example, methods for genetically
25 engineering bacteria are described in U.S. Patent
No. 4,259,444 to Chakrabarty, U.S. Patent No.
4,278,765 to Debabov et al., U.S. Patent No.
4,294,927 to Alfoldi et al., U.S. Patent
No. 4,237,224 to Cohen et al., U.S. 4,332,892 to
30 Ptashne et al. and U.S. Patent No. 4,338,397 to
Gilbert et al. U.S. Patent No. 4,318,987 to
Murillo Araujo et al. and Ilgen, Christine; Farabaugh,
P. J.; Hinnen, A.; Walsh, Jean M.; and Fink, G. R.
In: Genetic Engineering, Vol. 1, eds. Jane K. Setlow
35 and Alexander Hollaender (New York, Plenum Press)
1979, pp. 117-132, describe genetic engineering

41 12 11 90

0108580

-4-

systems for fungi and yeasts. Methods for genetically engineering in animal cells have also been described. For review see Lloyd H. Graf, Jr., "Gene Transformation," American Scientist, vol. 70, pp. 496-505, Sept.-Oct. 1982.

5 Presently, there are no systems for genetic engineering in algae. There are no vectors, transformation mechanisms, or selection systems available. Accordingly, there is need for a genetic engineering
10 system where foreign DNA is first transferred to, and then expressed in, algae.

Agrobacterium are a group of plant pathogens that infect a number of dicotyledonous plants. These include sunflower, tobacco, and carrot. The bacteria
15 cannot infect monocotyledonous plants, such as grasses, orchids, or lilies. During the infection process, a small portion of the bacterial plasmid DNA, the T-DNA, is transferred to the plant cell and is incorporated into the plant genome. Chilton, Mary-
20 Dell, et al., "Stable Incorporation of Plasmid DNA into Higher Plant Cells: The Molecular Basis of Crown Gall Tumorigenesis," Cell, vol. 11, pp. 263-271, June 1977. The T-DNA is replicated and expressed causing the plant cell to have a completely altered
25 metabolism. In addition the infected plant cell synthesizes a variety of anomalous amino acids or opines.

 Since DNA is transferred from the bacterial cell to the plant cell, many groups have considered
30 using crown gall bacteria (Agrobacterium tumefaciens), a species of Agrobacterium, as a vector to genetically engineer plant cells. In one scheme, a desired gene is attached to the T-DNA region of the Ti plasmid by gene splicing techniques. The crown gall
35 bacterial strain would be one that can infect and transfer DNA to the plant cell but would not cause

4 12 11 90

0108580

-5-

the plant cell to become tumorous. The plant cell is then infected with the crown gall bacteria, and the foreign gene is transferred to the plant cell with the T-DNA. The result is a genetically engineered plant cell that contains the foreign gene.

Several groups have been able to transfer foreign genes to plant cells using crown gall bacteria. However, many of these foreign genes have not been expressed by the plant cells. See Garfinkel, David J., et al. "Genetic Analysis of Crown Gall: Fine Structure Map of the T-DNA by Site-Directed Mutagenesis," Cell, vol. 27, pp. 143-153, Nov. 1981. Crown gall bacteria and other species of Agrobacterium, however, have not been used to transfer foreign genes into algal cells. As indicated above, there are presently no systems available for the genetic engineering of algae.

Therefore, it is an object of this invention to create a method for genetically engineering algae in which foreign genes may be transferred into algal cells, replicated, and expressed. It is a further object of this invention to create a genetically engineered algal cell in which foreign DNA is expressed.

The objects of this invention have been attained through a method of introducing foreign genes into green algae utilizing the plasmid DNA of Agrobacterium as a vector. Specifically, foreign genes composed of either eukaryotic or prokaryotic DNA can be transferred into host green algal cells utilizing Agrobacterium T-DNA as a vector wherein the foreign genes are incorporated into the algal genome, and replicated and expressed by the algal biosynthetic machinery. The method can be used to produce a genetically engineered green algal cell in which foreign DNA is incorporated, replicated and expressed.

4 12 11 90

0108580

-6-

I have found that Agrobacterium is able to infect green algae in a process similar to the process in which it infects higher plants. This is surprising in view of the previously recognized host range of Agrobacterium which excluded all lower plants. See, for example, De Cleene, Marcel and De Lev, Jozef, "The Host Range of Crown Gall" The Botanical Review, vol. 42(4), 1976, pp. 389-466. It is also surprising that green algae are infected considering Agrobacterium is apparently unable to infect blue-green algae as is evidenced in the Examples below. One would not expect from the limited documented host range of Agrobacterium and its inability to infect blue-green algae, that the bacteria would infect green algae. However, I have found that Agrobacterium is able to infect green algae and transfer bacterial T-DNA into the algal genome.

The first step of the invention requires the incorporation of foreign DNA into the vector. Because it is the T-DNA section of the Agrobacterium plasmid DNA which is incorporated into the genome of the infected cell, it is essential that the gene or genes to be transferred are incorporated within the T-DNA sequence. The desired foreign gene or genes can be inserted into the T-DNA using gene splicing procedures known in the art. One such procedure is described in Garfinkel, David J., et al., "Genetic Analysis of Crown Gall: Five Structure Map of the T-DNA by Site-Directed Mutagenesis," Cell, vol. 27, pp. 143-153, November, 1981, which is incorporated by reference.

The foreign genes which are inserted into the DNA can be composed of either eukaryotic or prokaryotic DNA. The biosynthetic machinery of green algae is able to recognize promoter sequences and express

44 12 11 90

0108580

-7-

genes from both systems unlike green plants which recognize only eukaryotic promoters.

A wide variety of genes can be employed as the foreign genes from a wide variety of sources. Any
5 intact gene can be employed which can be bonded to the plasmid vehicle.

One group of proteins which are directly useful are hormones. Illustrative hormones include para-
thyroid hormone, growth hormone, gonadotrophins,
10 (FSH, luteinizing hormone, chorionogonadotrophin, and glycoproteins), insulin, ACTH, somatostatin, prolactin, placental lactogen, melanocyte stimulating hormone, thyrotropin, parathyroid hormone, calcitonin, enkephalin, and angiotensin.

Other proteins of interest include serum proteins, fibrinogen, prothrombin, thromboplastin, globulin,
e.g., gamma-globulins or antibodies, heparin, antihemo-
15 philia protein, oxytocin, albumins, actin, myosin, hemoglobin, ferritin, cytochrome, myoglobin, lacto-
20 globulin, histones, avidin, thyroglobulin, interferin, kinins and transcortin.

Where the gene or genes produce one or more enzymes, the enzymes may be used for fulfilling a wide variety of functions. Included in these functions
25 are nitrogen fixation, production of amino acids, e.g., polyiodothyronine, particularly thyroxine, vitamins, both water and fat soluble vitamins, anti-microbial drugs, chemotherapeutic agents, e.g., anti-tumor drugs, polypeptides and proteins e.g. enzymes
30 from apoenzymes and hormones from prohormones, diagnostic reagents, energy producing combinations, e.g., photosynthesis and hydrogen production, prostaglandins, steroids, cardiac glycosides, coenzymes, and the like.

35 The enzymes can be individually useful as agents separate from the algal cell for commercial applica-

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0108580

-8-

tions, e.g., in detergents, synthetic transformations, diagnostic agents and the like. Enzymes are classified by the I.U.B. under the classifications as I. Oxidoreductases; II. Transferases; III. Hydrolases; 5 IV. Lyases; V. Isomerases; and VI. Ligases.

It is preferable when inserting foreign genes into the vector that a selection marker also be inserted. A selection marker is a gene or set of genes which codes for a characteristic which allows 10 for separation of cells which have been successfully genetically engineered from those which have not. Selection markers are well-known in the art and include sequences which confer antibiotic resistance, sequences which detoxify and sequences restoring 15 prototrophy. The selection system to be used in a particular system will be dependent upon several factors including, but not limited to, the algal strain used and the desired end product.

Once the vector has been prepared, algae are 20 infected by the Agrobacterium strain containing the engineered plasmid. It has been found in plant cells that metabolically active bacterial cells produce more tumors. Therefore, it is preferable to use Agrobacterium cultures which are in the mid- 25 to late-log phase. Cells in the cultures should be washed with sterile algal growth medium to remove bacterial growth medium in order to prevent bacteria from overgrowing the algae when algal cells are plated on selective medium. In plants, the cells 30 must be actively dividing in order to be infected by the bacteria. It is therefore preferable to use algal cells which are in the actively dividing mode or mid- to late-log phase. Cultures of the bacteria and algae are mixed together on algal growth medium. 35 Cells are allowed to incubate for 30 seconds to 15 minutes, preferably 5 minutes, at less than 32°C.

4 12 11 90

0108580

-9-

Agrobacterium are not known to infect plants at temperatures of 32°C or higher. The actual temperature for infection is preferably the optimum temperature for growth of the particular algal species employed. Significantly lower temperatures would not allow the algal cells to actively divide, a necessary condition for infection. During infection the T-DNA of the bacterial plasmid is incorporated into the algal DNA where it is replicated and expressed.

Agrobacterium can be used as a vector in the engineering of all green algae. Green alga include the members of the chlorophyta phylum. This phylum includes, for example, the genera Chlamydomonas, Dunaliella, and Chlorella. This phylum particularly includes Chlamydomonas reinhardtii, Protosiphon botryoides, and Chlorella pyrenoidosa.

Currently, several species of Agrobacterium have been identified which can be used in this process. Species which would be preferable due to extensive information which has been compiled regarding the genetic compositions of their plasmids and their infection processes include Agrobacterium rhizogenes and Agrobacterium tumefaciens.

EXAMPLES

In examples listed below, the following cultures and conditions were used.

The green alga Chlamydomonas reinhardtii (UTEX 2246) was grown on CA medium (Table 1).

The green alga Protosiphon botryoides (Cambridge Culture Collection 731/1a) was grown on 625 medium (Table 1).

The blue-green alga Anacystis nidulans (ATCC 27344) was grown on 625 medium (Table 1).

The 320 strain of Agrobacterium tumefaciens was obtained from Drs. Eugene Nester and Milton Gordon

11 12 11 90

0108580

- 10 -

(Garfinkel et al., 1981) and was grown on nutrient medium.

The aforementioned strains are all available to the public strains UTEX 2246, CCC 731/1A and ATCC 27344 having been made available through the respective depositories. The 320 strain may be obtained from Drs Eugene Nester and Milton Gordon, Department of Biochemistry, University of Washington, Seattle, Washington 98195.

The 320 strain contains the transposon Tn5 in the T-DNA region of the Ti plasmid. Tn5 is a prokaryotic gene that codes for the protein aminoglycoside phosphotransferase (3')-II (APH). This protein detoxifies the antibiotics neomycin, kanamycin, and G418 by transferring the gamma phosphosphate group of ATP to the antibiotic. These antibiotics are toxic to many different species. Some bacteria and yeast cells which contain the Tn5 gene are able to grow in the presence of the antibiotics.

The aqueous extract from agar was prepared by stirring 10 g of Difco-Bacto agar in 1 litre of water for 2 hours. The suspension was filtered through 2 layers of Miracloth. One liter of CA agar medium was made using 1 liter aqueous extract.

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-11-

Table 1
Algae Growth Media
CA Medium

		<u>mg/l</u>
5	NH ₄ Cl	50
	MgSO ₄ 7H ₂ O	20
	CaCl ₂ 2H ₂ O	10
	K ₂ HPO ₄	720
	KH ₂ PO ₄	360
10	NaAcetate	2000

Plus 1 ml of Hutners trace elements per liter
medium:

		<u>g/l</u>
15	Na ₂ EDTA	50
	H ₃ BO ₃	11.4
	ZnSO ₄ 7H ₂ O	22
	MnCl ₂ 4H ₂ O	5.1
	FeSO ₄ 7H ₂ O	5
20	CoCl ₂ 6H ₂ O	1.6
	CuSO ₄ 5H ₂ O	1.6
	(NH ₄) ₆ Mo ₇ O ₂₄ 4H ₂ O	1.1

625 Medium

		<u>mg/l</u>
25	NaNO ₃	496
	K ₂ HPO ₄	39
	MgSO ₄ 7H ₂ O	75
	CaCl ₂ 2H ₂ O	36
30	Fe citrate	6
	Na ₂ SiO ₃ 9H ₂ O	58
	Na ₂ CO ₃	20
	Citric acid	6
	Na ₂ EDTA	1

35

Ph 7.5

4 12 11 90

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-12-

EXAMPLE I

A 50 ml. culture of the 320 strain was grown to mid- to late-log phase. The cells were collected by centrifugation and washed twice with sterile CA medium. The cells were resuspended in 5 ml. of medium. About 10^6 cells from 2-day-old Chlamydomonas cultures (mid- to late-log phase) were mixed with 0.2 ml. of the bacterial suspension at room temperature. The mixture was plated onto CA medium with 10 mg/l G418 sulfate (potency 500 mcg/mg), 25 mg/l streptomycin, and 25 mg/l kanamycin.

The plates were incubated at 27°C. 27°C was chosen because it is the optimum temperature for growth of Chlamydomonas. Untreated cells from the alga Chlamydomonas reinhardtii were killed when grown in the presence of the antibiotic G418. Algal colonies composed of the algal cells mixed with the Agrobacterium strain became visible within 4-5 days and grew vigorously in the presence of the antibiotic. The results indicate that the 320 strain infected and transferred the Tn5 gene to the algae and the algae were able to express the Tn5 gene to make a functional protein. Colonies of the Agrobacterium infected Chlamydomonas were isolated and grown on CA agar medium containing 10 mg/l G418 and 10 mg/l carbenicillin.

The infected Chlamydomonas cells also showed other indications of altered metabolism. Whereas normal Chlamydomonas cells grow equally well on liquid or solid media, infected cells would grow only in a liquid medium that had been supplemented with an aqueous extract of the Difco-Bacto agar. Cells, which normally grow as single cells, grew as a clumped mass. Alterations in morphology were

M 12.11.90

0108580

-13-

attributed either to the expressions of other regions of the T-DNA or the disruption of a normal algal gene or genes due to insertion of the T-DNA.

To further substantiate this evidence, several tests were performed on the infected cells. Proteins were isolated and assayed from cultures of infected Chlamydomonas cells from which crown gall bacterial cells had been removed. Results show the presence of the functional APH protein which confers antibiotic resistance indicating that the Tn5 gene had been transferred and expressed. Protein assays also indicated the presence in the algal cells of octopine synthase. Octopine synthase is protein which is coded for by the T-DNA and which contains a eukaryotic promoter.

Analysis of the APH and the octopine synthase proteins by SDS polyacrylamide gel electrophoresis revealed that these proteins were properly synthesized by the algal biosynthetic system. The infected Chlamydomonas cells were able to synthesize faithfully two genes of bacterial origin: the Tn5 gene and the octopine synthase gene. The Tn5 gene contains a bacterial promoter, and the octopine synthase gene presumably contains a plant promoter, since the gene is expressed only in Agrobacterium infected plant cells. The Chlamydomonas cells were able to recognize both types of promoters. This characteristic makes Chlamydomonas a very desirable host for gene transfer experiments.

DNA hybridization assays were also performed to ascertain incorporations of the Agrobacterium DNA into the algal DNA. Results indicate the presence of T-DNA in the infected Chlamydomonas cell DNA.

EXAMPLE II

As in Example 1, a 50 ml. culture of the 320 strain was grown to mid to late log phase. The

-14-

cells were collected by centrifugation and washed twice with sterile 625 media. The cells were re-suspended in 5 ml. of media.

For the infection of Protosiphon, about 10^6 cells from 4-6-day-old cultures (mid- to late-log phase) were mixed with 0.2 ml. of the bacterial suspension. The mixture was incubated at room temperature for five minutes and was then plated onto 625 medium supplemented with 100 mg/l kanamycin and 25 mg/l streptomycin and allowed to grow at 27°C. Colonies became visible in about 7-10 days.

Kanamycin is toxic to the Protosiphon cells, and the untreated algal cells did not grow on the selective medium. The infected algal cells grew in the presence of the antibiotic. These results indicate that the 320 crown gall bacterial strain infected and transferred the Tn5 gene to Protosiphon and that the algal cells were able to express the Tn5 gene to make a functional protein.

20 EXAMPLE III

As in Example I, a 50 ml. culture of the 320 strain was grown to mid- to late-log phase. The cells were collected by centrifugation and washed twice with sterile 625 media. The cells were resus-
25 pended in 5 ml. of media.

For the infection of Anacystis, about 10^6 cells from 2-3-day-old cultures (mid- to late-log phase) were mixed with 0.2 ml. of the bacterial suspension. The mixture was incubated at room temperature for
30 five minutes and was then plated onto 625 media supplemented with 10 mg/l kanamycin and allowed to grow at 27°C. No algal colonies were formed after four weeks.

These results indicate that Agrobacterium was
35 unable either to infect blue-green algae or transfer a functional gene into the algal cells.

44 12 11 90

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-15-

The above examples are presented for the purpose of illustration only and are not intended to be limitations of the invention. Additional uses of the methods and products will be apparent to those skilled in the art.

Thus for example strains of green algae and Agrobacterium sp. other than those specifically referred to in the examples may be employed.

CLAIMS:

1. A method for introducing foreign DNA into green algae comprising: infecting a green algal cell with genetically altered Agrobacterium which contains a biologically functional foreign gene incorporated within the T-DNA sequence of the bacterial plasmid.
2. The method of Claim 1 wherein the Agrobacterium is selected from Agrobacterium tumefaciens and Agrobacterium rhizogenes.
3. The method of Claim 1 or Claim 2 wherein the foreign DNA is selected from eukaryotic DNA, prokaryotic DNA and combinations thereof.
4. The method of any preceding claim wherein the green algae comprises algae chosen from the species Chlamydomonas reinhardtii, Protosiphon botryoides, and Chlorella pyrenoidosa.
5. The method of any preceding claim wherein the foreign DNA comprises a selection marker.
6. A genetically engineered green algal cell containing the T-DNA sequence of an Agrobacterium plasmid wherein said T-DNA contains a biologically functional foreign gene.
7. A genetically engineered green algal cell according to Claim 6 wherein the Agrobacterium is selected from the group consisting of Agrobacterium tumefaciens and Agrobacterium rhizogenes.
8. A genetically engineered green algal cell according to Claim 6 or Claim 7 wherein the foreign DNA is selected from the group consisting of eukaryotic DNA, prokaryotic DNA and combinations thereof.
9. The composition of matter of Claim 11 wherein the green algae comprises algae chosen from the species Chlamydomonas reinhardtii, Protosiphon botryoides, and Chlorella pyrenoidosa.

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-17-

10. A genetically engineered green algal cell according to any of Claims 7 to 9 wherein the foreign DNA comprises a selection marker.

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-18-

0108580

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Dear Sirs

European Patent Application No. 83306603.8 ✓
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Permission is requested to correct the following errors occurring in
Claim 9 of the above application :-

~~Claim 9 line 1 replace "Claim 11" by "Claim 6"~~
Claim 9 line 3 - replace "botoryoides" by "botryoides"

Regarding the second of these proposed amendments, it will be noted
that the species name botryoides is correctly spelled in Claim 4 and
in line 17 on page 9.

Replacement copies of page 16 are filed herewith in triplicate.

Yours faithfully

Stephen David Ritter ✓
STEPHEN DAVID RITTER
MATHYS & SQUIRE

The request for correction is allowed under
R. 88 EPC / with the exception of the deleted
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EUROPEAN SEARCH REPORT

Application number

DOCUMENTS CONSIDERED TO BE RELEVANT			EP 83306603.8
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 7)
A	FR - A - 2 438 087 (SNAM PROGETTI S.P.A.) * Page 3, line 21 - page 4, line 36 *	1	C 12 N 15/00 C 12 N 1/12 C 12 N 1/20 C 12 N 5/00
A	DE - B - 1 211 017 (KABUSHIKI KAISHA YAKULT HONSHA) * Claim *	1, 4, 9	
A, D	US - A - 4 259 444 (CHAKRABARTY) * Abstract *	1	
A, D	US - A - 4 237 224 (COHEN et al.) * Abstract *	1	
			TECHNICAL FIELDS SEARCHED (Int. Cl. 7)
			C 12 N
The present search report has been drawn up for all claims			
Place of search VIENNA		Date of completion of the search 02-02-1984	Examiner WOLF
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